

FORM PTO-1449(Modified) LIST OF PATENTS AND PUBLICATIONS FOR APPLICANT'S INFORMATION DISCLOSURE STATEMENT	ATTY. DOCKET NO.: V0179/7001	SERIAL NO.: 09/485005
	APPLICANT: Wanker et al.	
	FILING DATE: Herewith	GROUP: Not yet assigned

U.S. PATENT DOCUMENTS

Exam Init	Ref Des	Document No.	Date	Name	Class	Sub Class	FILING DATE If Appropriate
919		5,234,814	08/10/93	Card et al.	435	7.21	06/01/89
919		5,723,301	03/03/98	Burke et al.	435	7.1	11/03/95

FOREIGN PATENT DOCUMENTS

	Country & Doc. No. (11)	Pub. Date (43)	Class	Sub Class	Translation Yes No
919	EP 0 206302A2	30.12.86			✓
919	EP 0 293 249 A1	30.11.88			✓
919	EP 0 854 364 A1	22.07.98			✓
919	WO95/29243	02.11.95			✓
919	WO96/12544	02.05.96			✓
919	WO96/28471	19.09.96			✓
919	WO97/17445	15.05.97			✓

OTHER ART

(Including Author, Title, Date, Pertinent Pages, Publication, Etc.)

919		Kisilevsky et al., "Arresting amyloidosis <i>in vivo</i> using small-molecule anionic sulphonates or sulphates: implications for Alzheimer's disease", (2/95), pp. 143-148, <i>Nature Medicine</i> , Vol. 1, No. 2.
919		Trottler et al., "Polyglutamine expansion as a pathological epitope in Huntington's disease and four dominant cerebellar ataxias", (11/23/85), pp. 403-406, <i>Nature</i> Vol. 378.
919		Gutekunst et al., "Identification and localization of huntingtin in brain and human lymphoblastoid cell lines with anti-fusion protein antibodies", (9/95), pp. 8710-8714, <i>Pro. Natl. Acad. Sci. USA</i> Vol. 92 Neurobiology.
919		M.F. Perutz, "Glutamine repeats and inherited neurodegenerative diseases: molecular aspects", (1996) pp. 848-858, <i>MRC Laboratory of Molecular Biology</i> .

EXAMINER <i>Denise B. Sabel</i>	DATE CONSIDERED <i>10/24/01</i>
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EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant

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94		Mangiarini et al., "Exon 1 of the <i>HD</i> gene with an expanded CAG repeat is sufficient to cause a progressive neurological phenotype in transgenic mice", (11/01/96), pp. 493-506, <i>Cell</i> , Vol. 87.
94		Stott et al., "Incorporation of glutamine repeats makes protein oligomerize: implications for neurodegenerative diseases", (7/95), pp. 6509-6513, <i>Pro. Natl. Acad. Sci. USA</i> , Vol. 92, Biochemistry.
94		Merlini et al., "Interaction of the anthracycline 4'-iodo-4'-deoxydoxorubicin with amyloid fibrils: Inhibition of amyloidogenesis", (3/95), pp. 2959-2963, <i>Proc. Natl. Acad. Sci. USA</i> , Vol. 92, Medical Sciences.
94		Scherzinger, et al., "Huntingtin-encoded polyglutamine expansions form amyloid-like protein aggregates <i>in vitro</i> and <i>in vivo</i> ", (8/8/97), pp. 549-558, <i>Cell</i> , Vol. 90.
94		Davies et al., "Formation of neuronal intranuclear inclusions underlies the neurological dysfunction in mice transgenic for the HD mutation", (8/8/97), pp. 537-548, <i>Cell</i> , Vol. 90.
94		Derwent, "Nucleic acid fragments associated with spinocerebellar ataxia type 2-contain increased number of CAG repeat region compared to normal gene", (5/7/98), XP-002101183. <i>Abstract</i>
94		Tateishi et al., "Removal of causative agent of creutzfeldt-jakob disease (CJD) through membrane filtration method", (1993), pp. 357-362, <i>Membrane</i> , 18(6).

EXAMINER <i>Enileene A. Buhel</i>	DATE CONSIDERED <i>10/24/01</i>
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